

CC TO JUDGE KU

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7 UNITED STATES DISTRICT COURT BY
WESTERN DISTRICT OF WASHINGTON
8 AT SEATTLE

9 Q-PHARMA, INCORPORATED,

) No C01-1312P

10 Plaintiff-Counterdefendant,

11 v

12 THE ANDREW JERGENS
CORPORATION, INCORPORATED,

)
THE ANDREW JERGENS
COMPANY'S MOTION AND
MEMORANDUM OF POINTS AND
AUTHORITIES IN SUPPORT OF ITS
MOTION FOR SANCTIONS
PURSUANT TO FED. R. CIV. P. 11

13 Defendant-Counterclaimant

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NOTE FOR HEARING: JULY 12, 2002



ORIGINAL

THE ANDREW JERGENS COMPANY'S MOTION FOR
SANCTIONS PURSUANT TO RULE 11

Case No C01-1312P

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**THE ANDREW JERGENS COMPANY'S MOTION FOR
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I

INTRODUCTION

Q-Pharma and its attorneys have violated Fed R Civ P 11 Q-Pharma and its attorneys violated Rule 11 by filing the complaint in this suit when they knew that the '373 patent was invalid, and when they knew or should have known, had they conducted a reasonable prefilng investigation, that the claims of the '373 patent were invalid Q-Pharma's attorneys also violated Rule 11 by filing the complaint either without interpreting the claims of the '373 patent or interpreting them in a frivolous manner Q-Pharma and its attorneys violated Rule 11 again when they filed Q-Pharma's Reply, in which it denied these facts in total disregard of the evidence Q-Pharma and its attorneys should be sanctioned pursuant to Rule 11 in an amount at least as great as Jergens' attorney's fees and costs in this suit

II

FACTS

On August 24, 2001, Q-Pharma brought this suit against Jergens, alleging direct infringement, contributory infringement and active inducement [of others] to infringe the claims of U.S. Patent No. 4,654,373 ("the '373 patent") based solely on Jergens' manufacture, advertising and sale of Jergens' Curel® Age Defying Therapeutic Moisturizing lotion containing Coenzyme Q₁₀ (the "Accused Product")

On October 5, 2001, Jergens filed an Answer, Affirmative Defenses and Counterclaims, which included counterclaims for (I) declaratory judgment of invalidity of the '373 patent, (II) declaratory judgment of noninfringement of the '373 patent, (III) declaratory judgment of unenforceability of the '373 patent due to patent misuse, and (IV) antitrust violations under Section 2 of the Sherman Act.

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1 On January 17, 2002, Jergens filed a motion for summary judgment of invalidity of
2 the claims of the '373 patent, an opposition to which was filed on February 25, 2002, by Q-
3 Pharma

4 On April 30, 2002, Jergens filed for a motion for summary judgment of
5 noninfringement of any of the claims of the '373 patent In response, Q-Pharma filed a
6 Nonopposition on May 9, 2002, and sought a voluntary dismissal with prejudice of their
7 patent infringement claims

8 On May 16, 2002, Q-Pharma filed an Undertaking Not to Sue Jergens for
9 infringement of the '343 patent with respect to its sale of the Accused Product

10 On May 17, 2002, a hearing was held by this Court on Jergens' motion for summary
11 judgment of invalidity of the claims of the '373 patent At that hearing, Jergens put Q-
12 Pharma on notice that its conduct in this suit violated Fed R Civ P 11

13 III

14 ARGUMENT

15 A Legal Standards for Rule 11

16 In relevant part, Federal Rule of Civil Procedure 11 states

17 By presenting to the court (whether by signing, filing, submitting, or later
18 advocating) a pleading, written motion, or other paper, an attorney or
19 unrepresented party is certifying that to the best of the person's knowledge,
information, and belief, formed after an inquiry reasonable under the
circumstances,--

20 (1) it is not being presented for any improper purpose, such as to
21 harass or to cause unnecessary delay or needless increase in the cost of
litigation,

22 (2) the claims, defenses, and other legal contentions therein are
23 warranted by existing law or by a nonfrivolous argument for the extension,
modification, or reversal of existing law or the establishment of new law,

24 (3) the allegations and other factual contentions have evidentiary
25 support or, if specifically so identified, are likely to have evidentiary support
after a reasonable opportunity for further investigation or discovery

26 Fed R Civ P 11(b) (emphasis added)

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1 The purpose of Rule 11 is to "deter baseless filings in the district courts and
 2 streamline the administration and procedure of the federal courts" Cooter & Gell v.
 3 Hartmax Corp., 496 U.S. 384, 393 (1990). Rule 11 imposes an affirmative duty among
 4 parties and their attorneys to reasonably investigate the factual bases contained in any paper
 5 filed. Id. "The rule requires litigants to 'stop and think' before initially making legal or
 6 factual contentions." Advisory Committee Note on Rule 11, 28 U.S.C.A., p. 260. More
 7 specifically, in a Complaint, Rule 11 imposes a duty to take the necessary care in its
 8 preparation in order to prevent an abuse of the judicial system because "[b]aseless filings put
 9 the machinery of justice in motion burdening courts and individuals alike with needless
 10 expense and delay." Cooter, 496 U.S. at 398.

11 The imposition of sanctions under Rule 11 does not require a showing of bad faith,
 12 but simply a showing of a lack of reasonable investigation Estate of Blas v. Winkler, 792
 13 F.2d 858, 860 (9th Cir. 1986). The reasonableness of an inquiry prior to filing is determined
 14 by applying an objective standard as to both the factual and legal components of the
 15 filing. See Business Guides, Inc. v. Chromatic Comms. Enters., Inc., 498 U.S. 533, 548-50
 16 (1991).

17 In a patent infringement context, a plaintiff bears a significant pre-filing investigation
 18 burden before asserting a patent claim, and that burden cannot be fulfilled by merely filing
 19 suit on a suspicion of infringement and then asking for discovery to prove the suspicions.
 20 See Refac Int'l, Ltd. v. Hitachi, Ltd., 921 F.2d 1247, (Fed. Cir. 1990), Micro Motion, Inc. v.
 21 Kane Steel Co., 894 F.2d 1318 (Fed. Cir. 1990).

22 B Q-Pharma Should Be Sanctioned for Its Refusal to Amend Its Reply

23 Rule 11 requires that denials of factual contentions in papers filed with the Court be
 24 warranted on the evidence. Fed. R. Civ. P. 11(b)(4). Q-Pharma's Reply denies several
 25 allegations, discussed in the following paragraphs, which denials are not warranted based on
 26 the evidence. Q-Pharma was put on notice that these denials were not warranted based on

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1 the evidence at the hearing held in this Court on May 17, 2002, and was provided with
 2 further notice by Jergens' service of this Motion for Sanctions Pursuant to Fed R Civ P 11
 3 on Q-Pharma 1 Q-Pharma has not amended its Reply despite having received notice that the
 4 denials contained therein do not have a sufficient evidentiary basis Accordingly, Q-
 5 Pharma's refusal to amend its Reply serves as a basis for the award of sanctions pursuant to
 6 Rule 11

7 Paragraph 16 of Jergens' Answer alleges that

8 Q-Pharma knows, and/or in the exercise of a reasonable pre-filing
 9 investigation should have known, that the amount of CoQ₁₀ in the Jergens'
 10 Curel® Lotion is below the ranges described in the '373 patent for both
 therapeutic and cosmetic purposes, and that to the extent Plaintiff Q-Pharma is
 asserting the product can be used for purposes proscribed by the '373 patent,
 CoQ₁₀ is not the principal active agent in such use

11 Q-Pharma denied these allegations in paragraph 2 of its Reply However, as discussed in
 12 further detail below, Q-Pharma could have run two simple tests to determine the amount of
 13 CoQ₁₀ present in the Jergens Curel® Lotion and to determine what constitutes the principal
 14 active ingredient of the Jergens Curel® Lotion, but failed to do so Furthermore, Q-Pharma
 15 could have determined that CoQ₁₀ was not the principal active ingredient in Jergens Curel®
 16 Lotion by simply reading the list of ingredients on the label Thus, Q-Pharma's denial of
 17 these allegations is not warranted based on the evidence

18 Paragraph 31 of Jergens' Answer seeks a declaration of invalidity of the '373 patent
 19 Paragraph 31 of Q-Pharma's Reply denies that Jergens is entitled to such a declaration
 20 However, as discussed in further detail below, Q-Pharma was made aware of the invalidity of
 21 the '373 patent in a letter from an accused infringer dated December 29, 2000 This letter
 22 clearly and undisputedly demonstrates the invalidity of the '373 patent Therefore, Q-
 23

24
 25 1 Jergens' Motion was served on Q-Pharma but not filed with the Court for 21 days pursuant
 26 to Rule 11(c)(1)(a)

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1 Pharma's denial that Jergens is entitled to a declaration of invalidity of the '373 patent is not
 2 warranted based on the evidence

3 Paragraph 47 alleges that bringing this suit violates Section 2 of the Sherman Act Q-
 4 Pharma denies this allegation in paragraph 34 of its Reply However, Q-Pharma brought this
 5 suit with the knowledge that the '373 patent was invalid and not infringed as discussed above
 6 Bringing a patent infringement suit with such knowledge is undisputably a violation of
 7 Section 2 of the Sherman Act PREI, Inc v Columbia Pictures Indus, Inc, 508 U S 49
 8 (Glaverbel Societe Anonyme v Northlake Marketing & Supply, Inc, 45 F 3d 1550
 9 (Fed Cir 1995), Walker Process Equip., Inc v Food Mach & Chem Corp, 382 U S 172
 10 (1965) Furthermore, Q-Pharma has as much as admitted that its primary reason for bringing
 11 this suit was to protect its licensees, including Alberto-Culver and Beiersdorf, rather than a
 12 belief that the '373 patent was infringed See Ex Y (Q-Pharma's draft letter to the Federal
 13 Trade Commission) at 1-2 Accordingly, Q-Pharma's denial that it has violated Section 2 of
 14 the Sherman Act was not warranted based on the evidence

15 Paragraph 37 of Jergens' Answer seeks a declaration of unenforceability due to
 16 misuse of the '373 patent Paragraph 24 of Q-Pharma's answer denies that Jergens is entitled
 17 to such a declaration However, as discussed above, the evidence shows that Q-Pharma's
 18 activities constitute anti-trust violations Therefore, these activities also constitute patent
 19 misuse Hunter-Douglas, Inc v Comforortex Corp, 44 F Supp 2d 145, 156 (S D N Y 1999)
 20 ("[c]learly, a patentee who uses a patent to violate the antitrust laws is guilty of patent
 21 misuse") Indeed, even activities that do not rise to the level of antitrust violations may
 22 constitute patent misuse Transparent-Wrap Mach Corp v Stokes & Smith Co, 329 U S
 23 637, 641 (1947), Virginia Panel Corp v MAC Panel Co, 133 F 3d 860, 872 (Fed Cir 1997)
 24 ("violation of section 2 of the Sherman Antitrust Act requires more exacting proof
 25 than suffices to demonstrate patent misuse") Therefore, Q-Pharma's denial that Jergens is
 26 not entitled to a declaration of patent misuse is not warranted based on the evidence

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1 As discussed above, Q-Pharma's denials of several allegations in Jergens' Answer are
 2 not warranted based on the evidence. Q-Pharma has failed to withdraw these unwarranted
 3 denials and therefore should be sanctioned pursuant to Rule 11.

4 **C Q-Pharma's Prefiling Claim Interpretation Was Frivolous**

5 Rule 11 requires an attorney who files a patent infringement suit to interpret the
 6 claims of the patent at issue before filing a complaint alleging patent infringement.
 7 Antonious v. Spalding and Evenflo Cos., Inc., 275 F.3d 1066, 1072 (Fed. Cir. 2002)
 8 "Because claim construction is a matter of law, an attorney's proposed claim construction is
 9 subject to the Rule 11(b)(2) requirement that all legal arguments be non-frivolous." Id.

10 Q-Pharma's prefiling interpretation of the "therapeutically effective amount"
 11 limitation of the claims, to the extent that any prefiling interpretation was attempted, is
 12 clearly frivolous. First, there is no evidence that Q-Pharma attempted to interpret any claim
 13 of the '373 patent prior to filing suit. No documents have been disclosed to Jergens that
 14 clearly indicate any prefiling claim interpretation was undertaken, nor does Q-Pharma's
 15 privilege log, attached hereto as Exhibit Z, refer to any claim charts or other documents that
 16 clearly indicate a prefiling interpretation was, in fact, performed.²

17 Second, to the extent that Q-Pharma's pleadings are reflective of a prefiling claim
 18 interpretation, it is clear that such an interpretation was frivolous. In the first substantive
 19 paper filed by Q-Pharma after the complaint, Q-Pharma took the position that

20 [T]estimony is made irrelevant by Jergens' admission that its Curel
 21 Therapeutic Moisturizing Lotion With Coenzyme Q₁₀ in fact contains CoQ₁₀.
 22 See Answer, Affirmative Defenses and Counterclaims ¶ 15 (admitting that
 23 accused product contains CoQ₁₀). Q-Pharma believes that this admission by
 24 Jergens establishes infringement, for it contends that any topical
 25 administration of CoQ₁₀ to treat skin, as would be done by a person using
 26 [Jergens'] Curel Therapeutic Moisturizing Lotion with Coenzyme Q₁₀,
 infringes the '373 patent.

2 Jergens will seek discovery on this issue to the extent permitted by the Court.

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1 Plaintiff's Memorandum in Opposition to Defendant's Motion to Transfer (October 22, 2001)
 2 at p 5, ll 16-18 (emphasis added) The position that any administration of CoQ₁₀, regardless
 3 of the amount of CoQ₁₀ present, infringes the '373 patent is plainly contradicted by the
 4 specification, which defines "therapeutically effective amount" by distinguishing between
 5 therapeutic and cosmetic uses, noting that for pharmacological purposes, CoQ₁₀ is present "as
 6 the active principle in amounts from 0.1 to 10%" See the '373 Patent at col 2, ll 53-60
 7 Thus, it is frivolous to take the position that any amount of CoQ₁₀ literally infringes the '373
 8 patent Furthermore, the file history clearly shows that the "therapeutically effective amount"
 9 limitation was added to the claims in order to overcome a prior art rejection See Jergens'
 10 Motion for Summary Judgment on Noninfringement at 3-5 Thus, taking the position prior to
 11 filing the Complaint that a topical administration of any amount of CoQ₁₀ would infringe the
 12 '373 patent under the doctrine of equivalents is also frivolous See Festo Corp v Shoketsu
 13 Kinzoku Kogyo Kabushiki Co., Ltd., 234 F 3d 558, 569 (Fed Cir 2000) (holding that the
 14 doctrine of equivalents is completely barred for a claim limitation that has been amended to
 15 overcome prior art) Thus, Q-Pharma's pre-filing interpretation that any topical administration
 16 of CoQ₁₀ infringes the '373 patent is frivolous and sanctionable under Rule 11

17 D Q-Pharma Should Have Known That the Accused Product Does Not Infringe the '373
 18 Patent

19 In addition to requiring an attorney to interpret the claim, Rule 11 also requires an
 20 attorney to make a reasonable pre-filing effort, independent of his client, to determine
 21 whether the accused product satisfies each of the claim limitations Antonious v Spalding
 22 and Evenflo Cos., Inc., 275 F 3d 1066, 1074 (Fed Cir 2002) Each claim of the '373 patent
 23 requires "a therapeutically effective amount" of Coenzyme Q₁₀ ("CoQ₁₀") and also requires
 24 that CoQ₁₀ be the "principal active ingredient" Q-Pharma's counsel failed to make a
 25 reasonable effort to determine whether the Accused Product met either of these limitations
 26 prior to filing this suit

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1 Q-Pharma failed to run either of two simple tests that would have confirmed
 2 noninfringement. The '373 patent defines a "therapeutically effective amount" of CoQ₁₀ to
 3 be at least 0.1%. The '373 patent, c 2, ll 53-54. Early in this suit, Q-Pharma admitted that
 4 the amount of CoQ₁₀ present in the Accused Product "could be resolved by any competent
 5 analytic chemist." Ex AA (Plaintiff's Memorandum in Opposition to Defendant's Motion to
 6 Transfer) at 6. Having a competent analytic chemist test the Accused Product would have
 7 revealed that the Accused Product contains far less than a "therapeutically effective amount."
 8 Ex BB (Maksimoski Decl.) ¶¶ 3, 7.

9 With regard to the "principal active ingredient limitation," Q-Pharma's expert, Mort
 10 Westman, testified that the "negative omission" test would determine which ingredient is the
 11 "principal active ingredient." Westman Depo at 38, l 3 to 41, l 15. When Jergens
 12 performed this very test, it clearly revealed that CoQ₁₀ was not the principal active ingredient
 13 as required by the claims. Ex CC (Witt Decl.) ¶ 8.

14 Q-Pharma could also have determined that the Accused Product does not infringe any
 15 claims of the '373 patent by simply reading the list of ingredients on the label. The Accused
 16 Product is a skin moisturizer. The label of the Accused Product reveals that it contains more
 17 glycerine than any ingredient other than water. Glycerine is a well-known skin moisturizer.
 18 Additionally, four other active ingredients are listed on the label before CoQ₁₀. Thus, by
 19 simply reading the label on the Accused Product, Q-Pharma knew or should have known that
 20 the CoQ₁₀ is not the "principal active ingredient" as recited in the claims.

21 Q-Pharma's failure to conduct either one of these two simple tests, either of which
 22 would have established noninfringement, is not reasonable. In Judin v. United States, the
 23 Federal Circuit reversed the trial court's decision not to sanction the patentee under Rule 11
 24 when the patentee "could have purchased a device for a minuscule amount, compared to the
 25 cost of the litigation" in order to determine whether the device infringed the patent. 110 F.3d
 26 780, 783 (Fed. Cir. 1997). Similarly, the aforementioned tests could have been performed

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1 for a minuscule amount, compared to the cost of this litigation Q-Pharma's additional
 2 failures to ask Jergens basic questions that could have established noninfringement, or read
 3 the ingredient list on the label of the Accused Product, are similarly not reasonable Thus, Q-
 4 Pharma failed to conduct an adequate prefilng investigation and therefore violated Rule 11
 5 by bringing this suit

6 In its Non-Opposition to Motion for Summary Judgment of Non-Infringement (the
 7 "Non-Opposition"), Q-Pharma attempted to establish a reasonable basis for bringing this suit
 8 by claiming that it believed the Accused Product had a therapeutically effective amount of
 9 CoQ₁₀ based on certain statements on the label of the Accused Product However, the
 10 Federal Circuit has made it clear that reliance on advertising as a basis for filing a suit is not
 11 sufficient In View Eng'g v Robotic Vision Sys, 208 F 3d 981 (Fed Cir 2000), the Federal
 12 Circuit noted that the patentee had not seen the alleged infringer's products prior to filing
 13 suit, but rather based its belief of infringement solely on the alleged infringer's "own
 14 advertising and its claims to customers as to what its machines did" Id. at 983 The court
 15 concluded that

16 [L]ittle inquiry, much less a reasonable one, was undertaken by [patentee's
 17 law firm] in the instant case Before filing []claims of patent infringement,
 18 Rule 11, we think, must be interpreted to require the law firm to, at a bare
 minimum, apply the claims of each and every patent that is being brought into
 the lawsuit to an accused device and conclude that there is a reasonable basis
 for a finding of infringement of at least one claim of each patent so asserted

19 Id. at 986

20 The similarity to this case is striking In its Non-Opposition, Q-Pharma admits that it
 21 simply assumed that the Accused Product infringed based on the advertising on the label

22 Based on the[] representations on the [Jergens Curel] label, Q-Pharma
assumed what any reasonable consumer reading the label would assume that
 23 Curel® Therapeutic Moisturizing Lotion with Coenzyme Q₁₀ contains enough
 24 CoQ₁₀ to accomplish the "therapeutic moisturizing" promised by its name, and
 thus had a therapeutically effective amount of CoQ₁₀ Q-pharma therefore
brought this action for infringement

25
 26 Q-Pharma's Non-Opposition at 2 (emphasis added)

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Relying on such an assumption as a basis for filing this patent infringement suit is simply not reasonable, especially when Q-Pharma admits that this assumption could have been confirmed by "any competent analytical chemist." Furthermore, this assumption does not provide Q-Pharma with a reasonable basis for a belief that CoQ₁₀ is the "principal active ingredient," which is also required by each claim of the '373 patent.

In addition to Q-Pharma's failure to run the aforementioned tests, Q-Pharma also failed to simply ask Jergens what amount of CoQ₁₀ is present in the Accused Product or whether CoQ₁₀ was the principal active ingredient in the Accused Product. The Federal Circuit has indicated that the failure to ask for information from an accused infringer, prior to suit, in order to verify infringement is a basis for Rule 11 sanctions. See Judin, 110 F.3d at 784 (imposing Rule 11 sanctions when patentee failed to ask for a sample product from the accused infringer so that infringement could be confirmed).

Q-Pharma's Non-Opposition also attempts to shift the blame for its lack of a reasonable basis to Jergens by complaining about Jergens' refusal to disclose the amount of CoQ₁₀ in the Accused Product in response to its discovery requests. Non-opposition at 2-3. However, these discovery requests came after this litigation was commenced. Rule 11 requires that the inquiry be taken before the suit is filed, not after. See Judin, 110 F.3d at 784. Accordingly, any post-filing refusal by Jergens to divulge the amount of CoQ₁₀ in the Accused Product does not excuse Q-Pharma from its obligation to make reasonable efforts prior to the filing of this suit to determine whether the Accused Product infringes the '373 patent.

Q-Pharma and their attorneys have made it clear that they felt free to pursue this reckless conduct because Q-Pharma is judgment-proof. See Ex DD (Letter from S. Dunwoody to S. Kelber dated May 20, 2002) at 1-2. This Court should not tolerate such

3 Jergens is aware that Federal Rule of Evidence 408 prohibits the use of settlement negotiations in compromising or attempting to compromise a claim as evidence to prove
(continued)

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1 behavior and should send a strong message to Q-Pharma and its attorneys that such reckless
 2 conduct will not be tolerated by sanctioning Q-Pharma and its attorneys pursuant to Rule 11

3 For the reasons discussed above, Q-Pharma has utterly failed to make a reasonable
 4 effort to determine whether the Accused Product infringes the '373 patent. Thus, Q-Pharma
 5 and its attorneys should be sanctioned under Rule 11 for bringing this suit.

6 E Q-Pharma Knew or Should Have Known That the '373 Patent Is Invalid

7 Bringing a patent infringement suit with the knowledge that the patent is invalid is a
 8 basis for awarding sanctions under Rule 11. Thermocycle Int'l, Inc v A F Henrichsen Sales
 9 Corp, 1991 U.S. Dist. LEXIS 8550, *7 (S.D.N.Y. 1991). Q-Pharma knew or should have
 10 known that the '373 patent is invalid. As fully briefed in Jergens' motion for summary
 11 judgment of invalidity of the '373 patent (the "Invalidity Motion"), the '373 patent is not
 12 entitled to the benefit of the first U.S. application (U.S. Ser. No. 476,556), but rather is only
 13 entitled to the filing date of the continuation-in-part application (U.S. Ser. No. 711,034).
 14 Invalidity Motion at 11-12. Because Q-Pharma is not entitled to the earlier filing date, the
 15 publication of its own Italian patent application corresponding to the earlier-filed U.S.
 16 application renders the '373 patent invalid. Id. at 17-21. Q-Pharma was put on notice of this
 17 issue when it received a letter dated December 29, 2000, from an accused infringer clearly
 18 showing that the '373 patent was not entitled to the filing date of the first U.S. application and
 19 was therefore invalid in light of a corresponding Belgian application (also based on the
 20 original Italian application). Ex. AA at 1-2. Thus, Q-Pharma either know or should have
 21 known that the '373 patent was invalid when it received this letter, which was approximately

22
 23 (continued)

24 liability for that claim. However, in this instance, Jergens offers Ex. DD in support of its
 25 motions for sanctions pursuant to Fed. R. Civ. P. 11, not to prove Q-Pharma's liability for any
 26 claim in this suit.

4 An attorney and client may be held jointly and severally liable for filings that are not well
 grounded in fact under Rule 11. Judin v. United States, 110 F.3d 780, 785 (Fed. Cir. 1997).

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1 seven months prior to the filing of the complaint in this suit⁵. Accordingly, Q-Pharma did
2 not have good-faith basis for bringing this infringement suit and Rule 11 sanctions are
3 appropriate

4 IV

5 CONCLUSION

6 For the foregoing reasons, Jergens respectfully request that this Court sanction Q-
7 Pharma pursuant to Rule 11 of the Federal Rules of Civil Procedure in an amount at least as
8 great as Jergens' attorney's fees and costs in defending itself in this suit.

9 RESPECTFULLY SUBMITTED this 21st day of June, 2002.

10 PIPER RUDNICK LLP

11 By 
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18 The Andrew Jergens Corporation, Inc.
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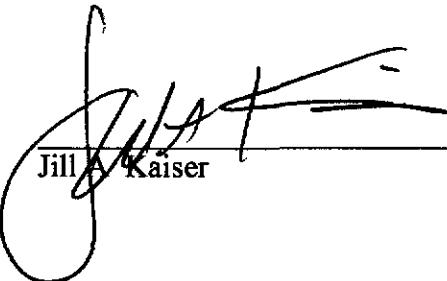
25 ⁵ Q-Pharma has not produced any opinion from counsel concerning this issue, nor does it
26 appear from their privilege log that any written opinion exists.

THE ANDREW JERGENS COMPANY'S MOTION FOR
SANCTIONS PURSUANT TO RULE 11 - 12
Case No. C01-1312P

1 **CERTIFICATE OF SERVICE**

2 I hereby certify that a true and correct copy of the above **THE ANDREW**
3 **JERGENS COMPANY'S MEMORANDUM OF POINTS AND AUTHORITIES IN**
4 **SUPPORT OF ITS MOTION FOR SANCTIONS PURSUANT TO FED. R. CIV. P. 11**
5 was served on June 21, 2002 via legal messenger on counsel of record at the following
6 address

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THE ANDREW JERGENS COMPANY'S MOTION FOR
SANCTIONS PURSUANT TO RULE 11 - 13
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